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Coronary Artery System Arising from a Solitary Coronary Ostium from the Right Sinus of Valsalva

Kristopher Pfirman^{1*}, Richard Pham1, Dominic Haddad², Connor Donley³, Mohammed Kazimuddin⁴ and Aniruddha Singh⁴

- ¹Department of Cardiology, The Medical Center at Bowling Green, Kentucky, USA
- ²Department of Osteopathic Medicine, University Of Pikeville, Kentucky, USA
- ³Department of Cardiology, University of Kentucky, Kentucky, USA
- ⁴Western Kentucky Heart and Lung, Kentucky, USA

Abstract

Coronary arteries arising from a single ostium are a rare anatomical variant. If an interarterial course is present this may manifest as angina and even sudden cardiac death. In our case, the left main coronary artery arises from a single ostium in the right sinus of Valsalva those courses in an inferior and posterior fashion. The anomalous left main coronary artery runs posterior to the aorta and courses along the left atrium. High risk features were absent. Due to the benign nature of the lesion, the patient was treated medically for coronary artery disease.

Introduction

In the general population, the blood supply to the heart is supplied by the right coronary artery and the left main coronary artery which arise from two separate coronary cusps. However, congenital variations to this anatomy can occur. In this case we present a patient with such coronary artery anomaly.

History of Presentation

A male in his fifties with obesity, obstructive sleep apnea compliant with continuous positive airway pressure at night, hypertension, uncontrolled diabetes, ongoing tobacco use, dyslipidemia and a history of cerebral vascular accident presented with chest tightness, epigastric pain, shortness of breath with exertion, and increased palpitations. Previous echocardiogram revealed a left ventricular ejection fraction of >55%. A 7-day Life watch monitor revealed sinus rhythm with no documented arrhythmias. Given the recent normal nuclear stress test performed at an outside facility and atypical yet ongoing symptoms, the patient underwent a diagnostic left heart catheterization.

Cardiac catheterization was done given the patient's high risk for Coronary Artery Disease (CAD) and chest pain complaints. The differential diagnosis would include occlusive coronary artery disease, Prinzmetal angina, microvascular coronary artery disease, coronary syndrome X, spontaneous coronary artery disease, coronary artery with interarterial course. The coronary angiogram revealed an anatomic anomaly of the left main coronary artery (Figure 1). There was no obstructive CAD found. Coronary Computed Tomography Angiography (CTA) was subsequently ordered to further assess the anatomy and origin of the coronary arteries. CTA revealed a single coronary ostium with common origin of the right and left coronary arteries arising from the right sinus of Valsalva. The left coronary artery coursing posteriorly between the aortic root and left atrium, without an abrupt acute angle, intramural course, or fish-mouth ovoid appearance of the ostium (Figure 2).

*Corresponding author: Kristopher Pfirman, Department of Cardiology, The Medical Center at Bowling Green, Kentucky, USA, E-mail: kpfirman19@gmail.com

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Figure 1: Coronary Angiography: LAO 30 degrees. Single coronary ostium arising from the right coronary cusp. Anomalous left main coronary artery arising from the right coronary artery.

Due to the mild atherosclerotic presence identified on coronary angiography, the course of action was to manage medically using beta blockade and calcium channel blockade in addition to aspirin and statin therapy. Given the inferior and posterior course of the anomalous left main coronary artery, conservative management was elected. Given the lack of the intra-arterial course that would prove a risk to the patient, surgery such as coronary reimplantation or coronary unroofing was deferred. The patient was then to be followed up in the outpatient clinic two weeks post discharge.

Discussion

Association with current guidelines/position papers/current practice: The left main coronary artery was found to have an anomalous origin in the right coronary cusp. Clinical presentation of patients with

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Figure 2: Coronary CTA images in the coronal, transverse, and sagittal planes revealing anomalous left main coronary artery from the right cusp of the sinus of Valsalva sharing a common ostium with the right coronary artery. The left main coronary artery takes aposterior and inferior course between the left atrium and aortic root.

coronary artery abnormalities range from asymptomatic to symptoms consistent with angina pectoris with possible sudden cardiac death, which is more common in young athletes [1]. It is estimated that up to 1.3% of the general population may have anomalous coronary artery origins [2]. Of this population, only 1.7% is estimated to have a single coronary artery arising from a solitary ostium (incidence of 0.024-0.066%) [2]. It has been shown in postmortem studies that patients with the anomalous left coronary artery arising from the right sinus, around 59% will die before age 20, and death usually occurs with vigorous exertion [3]. Whereas the incidence of sudden cardiac death related to any coronary artery anomaly in the general population under 40 years old is 0.6% [3]. Anatomical factors in this case, such as the left coronary artery not taking an interarterial or intramural course, absence of an acute angle, or fish-mouth ostium, makes this a more benign variant.

Treatment for these anomalies is dependent on the course of the anomalous vessel [4,5]. That's why classification of the anomaly is of the utmost importance, Lipton et al created the current classification system of single coronary arteries [2]. Coronary angiography is essential to establish the specific diagnosis [6]. Single coronary anomalies are usually asymptomatic, but an anomalous vessel which courses between the aorta and pulmonary trunk in an interarterial fashion is at high risk of compression, leading to anginal symptoms and possible sudden cardiac death [6,7]. These patients should receive referral to cardiothoracic surgery in order to bypass the vessel, reimplant the left coronary artery if able, or surgical unroofing of the vessel [8]. According to the 2018 AHA/ACC Guideline for the management of Adults with Congenital Heart Disease, surgery is a class one recommendation for symptoms or diagnostic evidence consistent with coronary ischemia attributable to the anomalous course of the coronary artery [9] There is a class IIa recommendation for anomalous origin of the left coronary artery from the right sinus in the absence of symptoms or ischemia or in the setting of ventricular arrhythmias [9].

Patients who have anomalous vessels which course around the great vessels and are not proceeding in an interarterial course are not at risk for compression and are therefore treated with conservative medical management as mentioned above. It should be noted that

professional associations have issued guidelines that state untreated carriers of anomalous origins of a coronary artery should not be involved in strenuous activities at risk for sudden cardiac death that exhibit a dangerous course [8]. Once treated, a re-evaluation should be conducted before exercise is resumed [8]. Coronary artery anomalies should also be accurately documented to reduce potential harm when patients receive care from other providers, such as undergoing aortic valve replacement [1,8].

Given the course of the left main coronary artery not running in interarterial fashion, the patient was elected to be treated medically for non-occlusive coronary artery disease. The patient was followed in the outpatient setting without any recurrence of his original presenting symptoms.

Conclusion

Anomalous coronary arteries are a rare finding which can be discovered on cardiac catheterization. Management is based upon symptomology as well as anatomical course. Our patient was found to have a variant that was not clinically significant for sudden cardiac death and therefore was managed with conservative medical management.

Conflict of Interest

No conflicts of interest to be presented.

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